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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/528,611	03/21/2005	Steven Gutteridge	BB1533USPCT	7575

7590 05/15/2007
E I du Pont de Nemours & Company
Legal Patents
Wilmington, DE 19898

EXAMINER

LI, RUIXIANG

ART UNIT	PAPER NUMBER
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1646

MAIL DATE	DELIVERY MODE
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05/15/2007

PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No. 10/528,611	Applicant(s) GUTTERIDGE ET AL.	
	Examiner Ruixiang Li	Art Unit 1646	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☐ Responsive to communication(s) filed on ____.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-33 is/are pending in the application.
- 4a) Of the above claim(s) ____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) ____ is/are allowed.
- 6) ☐ Claim(s) ____ is/are rejected.
- 7) ☐ Claim(s) ____ is/are objected to.
- 8) ☒ Claim(s) 1-33 are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on ____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
 Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
 Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. ____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- * See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|---|--|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413)
Paper No(s)/Mail Date. ____ |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08)
Paper No(s)/Mail Date ____ | 6) <input checked="" type="checkbox"/> Other: <u>Sequence alignment</u> . |

Election/Restrictions

1. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In accordance with 37 CFR 1.499, applicant is required, in response to this action, to elect a single invention to which the claims must be restricted.

- I. Claims 1-9, drawn to an isolated nucleotide fragment encoding a ryanodine receptor, a vector, and a host cell.
- II. Claim 10, drawn to a method to isolate nucleic acid fragments encoding ryanodine receptors and related polypeptides.
- III. Claims 11-15, drawn to an isolated polypeptide having ryanodine receptor activity.
- IV. Claims 16-20, drawn to a method for evaluating at least one compound for its ability to modulate calcium homeostasis or a method for evaluating at least compound which modulates ryanodine receptor activity.
- V. Claims 21 and 24-33, drawn to an isolated nucleic acid fragment encoding an insect ion channel, a method for expressing an isolated nucleic acid fragment encoding a toxic insect ion channel, and a recombinant constructor.
- VI. Claims 22 and 23, drawn to a method for identifying a nucleic acid sequence encoding an insect ion channel.

2. The inventions listed as Groups I-VI do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The technical feature linking Groups I-VI appears to be an isolated nucleotide fragment encoding a ryanodine receptor. However, claims 1-5 and 7-9 are anticipated by Takeshima et al. (FEBS Letters 337:81-87, 1994). Takeshima et al. teach a ryanodine receptor (Fig. 2) and its encoding DNA sequence (page 82). Since the ryanodine receptor of Takeshima et al. shares a high degree of homology with the amino acid sequence of SEQ ID NO: 128 of the present invention (see attached sequence alignment), the complementary sequence of the DNA sequence of Takeshima et al. comprises the complement of (a) of claim 1. It is noted that claim 1, part (b), does not require the complement of (a) is a full complement of (a) of claim 1 (i.e., over its entire length). Takeshima et al. further teach a recombinant construct and a host cell comprising the DNA sequence (bottom of right column of page 81) and use of CHO cells as host cells (top of right column of page 81). Thus, the teachings of Takeshima et al. meet the limitations of claims 1-5 and 7-9.

Therefore, the technical feature linking the inventions of Groups I-VI does not constitute a special technical feature as defined by PCT Rule 13.2, as it does not define a contribution over the prior art.

3. The special technical features of Groups I-III are an isolated nucleotide fragment encoding a ryanodine receptor, a vector, and a host cell, a method to isolate nucleic acid fragments encoding ryanodine receptors and related polypeptides, and an

2. The inventions listed as Groups I-VI do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The technical feature linking Groups I-VI appears to be an isolated nucleotide fragment encoding a ryanodine receptor. However, claims 1-5 and 7-9 are anticipated by Takeshima et al. (FEBS Letters 337:81-87, 1994). Takeshima et al. teach a ryanodine receptor (Fig. 2) and its encoding DNA sequence (page 82). Since the ryanodine receptor of Takeshima et al. shares a high degree of homology with the amino acid sequence of SEQ ID NO: 128 of the present invention (see attached sequence alignment), the complementary sequence of the DNA sequence of Takeshima et al. comprises the complement of (a) of claim 1. It is noted that claim 1, part (b), does not require the complement of (a) is a full complement of (a) of claim 1 (i.e., over its entire length). Takeshima et al. further teach a recombinant construct and a host cell comprising the DNA sequence (bottom of right column of page 81) and use of CHO cells as host cells (top of right column of page 81). Thus, the teachings of Takeshima et al. meet the limitations of claims 1-5 and 7-9.

Therefore, the technical feature linking the inventions of Groups I-VI does not constitute a special technical feature as defined by PCT Rule 13.2, as it does not define a contribution over the prior art.

3. The special technical features of Groups I-III are an isolated nucleotide fragment encoding a ryanodine receptor, a vector, and a host cell; a method to isolate nucleic acid fragments encoding ryanodine receptors and related polypeptides; and an

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isolated polypeptide having ryanodine receptor activity, respectively. The special technical features of Groups IV-VI are a method for evaluating at least one compound for its ability to modulate calcium homeostasis or a method for evaluating at least compound which modulates ryanodine receptor activity; an isolated nucleic acid fragment encoding an insect ion channel, a method for expressing an isolated nucleic acid fragment encoding a toxic insect ion channel, and a recombinant constructor; and a method for identifying a nucleic acid sequence encoding an insect ion channel, respectively.

4. Accordingly, Groups I-VI are not so linked by the same or a corresponding special technical feature as to form a single general inventive concept. Thus, unity of invention is lacking and restriction is appropriate.
5. Furthermore, this application contains claims directed to the following amino acid/nucleic acid sequences and each amino acid/nucleic acid sequence represents an *additional* invention group:

- (i). amino acid sequences set forth in SEQ ID NOS: 2, 4, 6, 8, 128, 130, 144, 146, and 63-119,

- (ii). nucleic acid sequences set forth in SEQ ID NOS: 1, 3, 5, 7, 9, 127, 129, 143, and 145.

According to PCT rule 13.2 and to the guidelines in Section (f)(i)(B)(1) of Annex B of the PCT administrative Instructions, all alternatives of a Markush Group must have a common structure. The amino acid/nucleic acid sequences are not regarded as being of similar nature because the nucleic acid/amino acid sequences

do not appear to share a common structure.

Applicant is advised that a reply to this requirement must include an identification of an amino acid/ nucleic acid sequence that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election. The Examiner notes that this is not a species election requirement; rather it sets forth additional invention groups.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48 (b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48 (b) and by the fee required under 37 CFR 1.17 (I).

Advisory Information

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Ruixiang Li whose telephone number is (571) 272-0875. The examiner can normally be reached on Monday through Friday from 8:30 am to 5:00 pm. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Nickol, can be reached on (571) 272-0835. The fax number for the organization where this application or proceeding is assigned is (571) 273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, please contact the Electronic Business Center (EBC) at the toll-free phone number 866-217-9197.



Ruixiang Li, Ph.D.
Primary Examiner
May 11, 2007

RUIXIANG LI, PH.D.
PRIMARY EXAMINER

GenCore version 5.1.7
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QM protein - protein search, using sw model

Run on: April 14, 2006, 01:34:23 ; Search time 75 Seconds
(without alignment)

3848.671 Million cell updates/sec

Title: US-10-668-767-128_COPY_1_3000

Perfect score: 15748

Sequence: 1 MAAEAGSGSEQDDVSFLRTE.....RLADNNHDIWAKKKKELVT 3000

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: pir1.*

2: pir2.*

3: pir3.*

4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	12295	78.1	5126	2 840450	ryanodine receptor
2	6928	43.4	5107	2 T29144	partial CDS - Caen
3	6770.5	43.0	4969	2 A37113	ryanodine receptor
4	6769	43.0	4967	2 872469	ryanodine receptor
5	6534.5	41.5	5037	1 A54161	ryanodine-binding
6	6453.5	41.0	5035	1 A46646	ryanodine receptor
7	6442.5	40.9	5032	1 A35041	ryanodine receptor
8	6439	40.9	5037	2 B35041	ryanodine receptor
9	6438.5	40.9	4859	2 874173	ryanodine receptor
10	6423	40.8	4869	2 866572	ryanodine receptor
11	6419	40.8	4868	2 B54161	ryanodine-binding
12	6413	40.7	4872	2 827272	ryanodine receptor
13	538	3.4	163	2 I47214	probable brain rya
14	520	3.3	162	2 I47213	cardiac muscle rya
15	327	2.1	2783	2 T31431	inositol 1,4,5-tri
16	323	2.1	2701	2 817796	inositol-trisphosp
17	296.5	1.9	2693	2 A40743	IP3 receptor, XIP3
18	292.5	1.9	2695	2 854974	type 1 inositol 1,
19	290.5	1.8	2670	2 A46719	inositol 1,4,5-tri
20	289	1.8	2713	2 A55713	inositol 1,4,5-tri
21	286.5	1.8	2734	2 B36579	inositol 1,4,5-tri
22	285	1.8	2671	2 A49873	inositol 1,4,5-tri
23	284	1.8	1966	2 T32552	hypothetical prote
24	284	1.8	2848	2 T32550	hypothetical prote
25	279	1.8	2749	1 ACMSIT	inositol 1,4,5-tri
26	279	1.8	2749	2 A36579	inositol 1,4,5-tri
27	277.5	1.8	2833	2 A43360	inositol 1,4,5-tri
28	205	1.3	1676	2 B71410	probable centromer
29	203.5	1.3	2541	2 T29340	hypothetical prote

30	198	1.3	3259	1 A56539	giantin - human
31	198	1.3	5369	2 T44807	myosubtilin synth
32	197.5	1.3	2954	2 T14156	kinesin-related pr
33	196.5	1.2	2218	2 B84683	hypothetical prote
34	192	1.2	1727	2 T50073	myosin-like coiled
35	192	1.2	3225	2 T52300	giantin - human
36	191	1.2	1882	2 T00069	hypothetical prote
37	189.5	1.2	2663	1 828261	centromere protein
38	188.5	1.2	1642	2 T08880	NMDA receptor-bind
39	188	1.2	4540	2 T30838	cytoplasmic dynein
40	182.5	1.2	2331	2 T25410	hypothetical prote
41	182.5	1.2	3187	2 JCS637	364K Golgi complex
42	180.5	1.1	2712	2 T05113	hypothetical prote
43	179	1.1	6669	2 S55024	nebulin, skeletal
44	178	1.1	4151	2 T13734	groovin gene prote
45	177	1.1	52	2 I46644	ryanodine receptor

ALIGNMENTS

RESULT 1

S40450

Ryanodine receptor/calcium release channel - fruit fly (Drosophila melanogaster)

C:Species: Drosophila melanogaster

C/Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 11-Jun-1999

C/Accession: S40450

R/Takeshima, H.; Nishi, M.; Iwabe, N.; Miyata, T.; Hosoya, T.; Masai, I.; Hotta, Y.

FEBS Lett. 337, 81-87, 1994

A/Title: Isolation and characterization of a gene for a ryanodine receptor/calcium rel

A/Reference number: S40450; MUID:94102409; PMID:8276118

A/Accession: S40450

A/Status: preliminary

A/Molecule type: DNA

A/Residues: 1-5126 <TR>

A/Cross-references: UNIPARC:UPI000017CF36

C/Suprafamily: ryanodine receptor; transcription initiation factor sigma region 1 homc

Query Match 78.1%; Score 12295; DB 2; Length 5126;

Best Local Similarity 77.5%; Pred. No. 0;

Matches 2348; Conservative 275; Mismatches 338; Indels 68; Gaps 22;

Qy	1	MAAEGSGSEQDDVSFLRTEDVCLSCATATGERSVCLAAEGFGNRHCFLENIADKHPDNL 60
Db	1	MAAEGG-SEQDDVSFLRTEDVCLSCATATGERSVCLAAEGFGNRHCFLENIADKHPDNL 59
Qy	61	SOCPVPIEQALSVEALQELVTAAGSTGKGTSGHRTLLYGNAILLRHNSDMYLACLST 120
Db	60	SOCPVPIEQALSVEALQELVTAAGSTGKGTSGHRTLLYGNAILLRHNSDMYLACLST 119
Qy	121	SSSQDELAFVGLQOHSQGEACWMTLHPASKORSGEKRVGDDLLVSVATERYLHTTK 180
Db	120	SSSNDKLSFDVGLQHSQGEACWMTVHPASKORSGEKRVGDDLLVSVATERYLHTTK 179
Qy	181	ENRGSIVNASSPHVTHMSVQPYGTGSRMKYGVYFGDVLRFPHGDECLTIPSTWTKG 240
Db	180	ENRGSIVNASSPHVTHMSVQPYGTGSRMKYGVYFGDVLRFPHGDECLTIPSTWTKG 239
Qy	241	QGNIVVYEGGSVMSQASLWLELARTKAGCFINWHPMRIRHITTTGRTYLVGNQENLY 300
Db	240	QGNIVVYEGGSVMSQASLWLELARTKAGCFINWHPMRIRHITTTGRTYLVGNQENLY 299
Qy	301	LVSRREANTASCAPLQEDDQKQVLEKDLVIGAPILIKYGVSTVIVQHSBTGLWSY 360
Db	300	LVYKGEASIAITTTFSM-QEKDDEKRVLEKDLVIGAPILIKYGVSTVIVQHSBTGLWSY 358
Qy	361	KSYRTKKGVGKVEKQAILHEEGNDCLQPSRQSEBSRTARVTRKCSLFTPTKINGL 420
Db	359	KSYRTKKGVGKVEKQAILHEEGNDCLQPSRQSEBSRTARVTRKCSLFTPTKIPAL 418
Qy	421	ETLQNRHSHPASVNLGEMVCLIEDLINTPAQPDENCSHEERKONKPAALNRQDLQES 480
Db	419	ETLQNRHSHPASVNLGEMVCLIEDLINTPAQPDENCSHEERKONKPAALNRQDLQES 478

Db 2612 YRLLEDAPLPDLATATILDESOGSESDALANRYIGNSILPLILKSKFTNEANYASL 2671
Qy 2672 LDATLTVYRLSKRMMLTKGOREAVSDFLVALTSAMQPSKMLKLLRLKLTVDVSKSEYTT 2731
Db 2672 LDATLTVYRLSKRMMLTKGOREAVSDFLVALTSAMQPSKMLKLLRLKLTVDVSKSEYTT 2731
Qy 2732 VALRLTLHYTERCAKYGSTGACGAGCASSDEKRLTMMLPSNIPOSLSKMDYRDELPG 2791
Db 2732 VALRLTLHYTERCAKYGSTGACGAGCASSDEKRLTMMLPSNIPOSLSKMDYRDELPG 2790
Qy 2792 KALPCLTAICGALPPDYSLGKXNDYDFYCKEQAQADLNPNPODQPIPTSSVALANDLMT 2851
Db 2791 KALPCLTAICGALPPDYSLGKXNDYDFYCKEQAQADLNPNPODQPIPTSSVALANDLMT 2848
Qy 2852 IVQKPSHHYDAWASRKENGWYVGGWSDSKTHPRLLKPNMNDYKERYKBPVREBL 2911
Db 2849 LVQKPSHHYDAWASRLSGWYVGGWSDSKTHPRLLKPNMNDYKERYKBPVREBL 2908
Qy 2912 KALLAICWSYHSEVDI PENNBSMRQSKGGRPEI--VTDSATPPDYNPHVDNML 2969
Db 2909 KGLLAIGWTYHSEVPLNHRGSTRQSK-----POINBFQNEGSPFNTPHVPDMSNL 2963
Qy 2970 TLRSEWONMARRLADNADINAKCKEHL 2998
Db 2964 TLRSEWONMARRLADNADINAKCKEHL 2992

RESULT 2
T19144
Partial CDS - Caenorhabditis elegans
C1: Species: Caenorhabditis elegans
C2: Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jul-2004
C3: Accession: T29144
R1: Payley, A.; Gattung, S.
A1: Description: The sequence of C. elegans cosmid K11C4.
A2: Reference number: 220577
A3: Accession: T29144
A4: Status: preliminary; translated from GB/EMBL/DBJ
A5: Molecule type: DNA
A6: Residues: 1-5107 (PAU)
A7: Cross-References: UNIPROT: Q94279; UNIPARC: UP1000011013D; EMBL: U64854; PIR: AAB18318.1
A8: Experimental notes: strain Bristol N2; clone K11C4
C1: Superfamily: ryanodine receptor; transcription initiation factor sigma region 1 homolog

Query Match 43.4%; Score 6028; DB 2; Length 5107;
Best Local Similarity 45.0%; Pred. No. 0;
Matches 1407; Conservative 519; Mismatches 850; Indels 350; Gaps 52;

Qy 113 MYTCLSTSSQDLAFVGLQKSGGACWTLHPASKORSSEKRVVGGDILLVSVAT 172
Db 1 MYTCLSTSSQDLAFVGLQKSGGACWTLHPASKORSSEKRVVGGDILLVSVAT 60
Qy 173 ERYLHATK---ENEVEI-----VNASPHVTHWSVQVGTGI 205
Db 61 ERYLHATK---ENEVEI-----VNASPHVTHWSVQVGTGI 120
Qy 206 SRMKYGVYTPGGDVLFPFGHDECLTIPSTWTKDGGQNVVYEGSGVMSQASLNLHLA 265
Db 121 MRYTRNGPLFENDVLALPHGDECLTIPENSEHPQNNVITYEGGAATVQASLNRVELI 180
Qy 266 RTWAGQPINWPHPRIRHTTCVRLGYNDQVELYVSRBATTCAPCLQEKDOKO 325
Db 181 RMQOIGALVGMEOPIRKHITSGRYGLVD--NSVLHYKKAQDFLTAPVMQNKDPKKQ 239
Qy 326 VLKDKOLEVIGAPIFYGDSTVIVQHSYGLWLSYKSYETKGGVGVKVEKQAILHEBK 385

Db 240 MLDKESBEGGNATIRYGETNAPIQHVKTQVLTSLSYQTTEVTKKGLGKVEKKAVALKQGH 299
Qy 386 MDDGLDLSRSQBSSESTARVIRKCSSLTPKINGLETLOQVRRHSHPFASVNLSEVMCL 445
Db 300 MDDCYTFMALESBSKARVIRKCSSVNLKIGIDALQLEBQNSQDTRVLDLNEVLKJM 359
Qy 446 EDLLNYPAQDEDEHEHKKONKFRALRRNODLPQEEGILNLILEADIKINVTISQGLAG 505
Db 360 EDLLNYPAQDEDEHEHKKONKFRALRRNODLPQEEGILNLILEADIKINVTISQGLAG 419
Qy 506 FLACDESGSQWEMISQYLYLLAALIKGHTNCAQFANSNRLNLTFSRLGSOASGEQTEM 565
Db 430 -LIGESTHVQKQSTYLLVLAAMIKNHYNCAQFASQRLDMLPRLSNPQASAB--GI 476
Qy 566 LDVLHCVLIDSPALNMWDEHIKVIISLEKCHGRDPKVLVDVLCSLCVGVNGVAVRSSQNN 625
Db 477 LDVLHCVLIDSPALNMWDEHIKVIISLEKCHGRDPKVLVDVLCSLCVGVNGVAVRSSQNN 536
Qy 626 ICDYLLPGRKLLLOTLVDHVSVRPNIPVGRVEGSAVYKVPYKVTMDHIEKTHMPH 685
Db 587 ITQYLLFGDOLLQTSMDHVSMMNVMGLWVEGSAVYKVPYKVTMDHIEKTHMPH 596
Qy 688 LRIGWANTTGVYVPGGEEKGNGVGDYLSYVFGAYLWSCGRKTPVNRTHABEVPVIR 745
Db 597 LRIGWANTTGVYVPGGEEKGNGVGDYLSYVFGAYLWSCGRKTPVNRTHABEVPVIR 745
Qy 746 KGVITGCLDLTVPIINPAPNGVRVTSFTNPNLEHPPVPPVISCSSKLSCLPFLGSEHGR 805
Db 652 KGVITGCLDLTVPIINPAPNGVRVTSFTNPNLEHPPVPPVISCSSKLSCLPFLGSEHGR 711
Qy 806 LRYAAPRGVSPVLESLLPOILSLSPCFYFGLNLSKRALAGPLVDD--TAPVTPVDTLQ 864
Db 712 LRYAAPRGVSPVLESLLPOILSLSPCFYFGLNLSKRALAGPLVDD--TAPVTPVDTLQ 864
Qy 865 ITLPTVYSRDKLAENTHEMAMKIEAGMYGQDREHLKHPCLVPFELPFAKRY 924
Db 770 TOLNHAETEMKQYAEHLHAWMKIELGWSYGETRSETSRKHPCLVTKFYLPSTKRY 829
Qy 925 DIQLAVOTLTKVLLQYIISLDKPPARVNVRL--PNEPFGMSNGYKCAPLDLSAVTLTPK 983
Db 830 NILLATTKYTHLTYHLITDPPCLRAVRLGPN--FOQNGYKFGPLDTHSIQLPAE 887
Qy 984 NDELVQDLAENTHAWRERIQGQTYGLNR--DSDHESPHLVYKVPKVDALKANRDT 1041
Db 888 LQPLTEALANTHNTWAKKIKRGWTFGLSEHVDATQKSPHLVYVQVDEIRKQANRES 947
Qy 1042 ASETVRLVYGMPLDPTGSHRALLLEASKOKOAFRTYRAEKNYAVSGKYVFFBI 1101
Db 948 ANEIRALQPLGLEPPAHR--DEYAEKELRAKONTYTRABATYAVCGKRYFFBI 1006
Qy 1102 LTAGPMRVGVNAHADNAPGMQDGNMVAFGYNEKRYSGMTESFGKQWAVGVGVPL 1161
Db 1007 LTAGPMRVGVNAHADNAPGMQDGNMVAFGYNEKRYSGMTESFGKQWAVGVGVPL 1065
Qy 1162 DLIDKTSISLCELLDALGSETTPADQ--GNFVPACTGLVGOKARLTGVQDVNTLKY 1220
Db 1086 DLNDRTISLNGELLDPGSENAFONVVVGGGLVPMTLGGGRLNFGQSSNLKP 1125
Qy 1221 FTTGCLQEGYFPVCMKRDVTHYTKQPIFNTYDEKIDTRI--DVTRIPAGSDTPCLK 1279
Db 1126 FTTGCLQEGYFPVCMKRDVTHYTKQPIFNTYDEKIDTRI--DVTRIPAGSDTPCLK 1185
Qy 1280 ISENTYET-----MEKANEPLRLSLPVI CNEFTIDEK-----ABRWELIKDQQLAKEA 1332
Db 1186 ILQKVTISBGPSPKAKBYIKLSLPVKNDTPVKNQKRTIRRLQKRYKPSVVSQI 1245
Qy 1333 VBAQPAHIDQIM--RSGFTMDNDIKGLHYDQNBELSPSRKQLPS-----RPFKSGMTRG 1387
Db 1246 RAGCIPEDDNEKEKGFILBSMLASKSHESDDESRSTNSKPSVOCDEPPA----- 1298
Qy 1388 VTIONTNTIQGVNGVGRSTSEBAQYKQYDGLGAGCLTPDDKKKGRGSPFFKPS-----K 1443
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Puentes, R., Sunar, M., Evans, A.D., McCaffery, A.R. and Windass, J.D.
Identification of a polymorphic ryanodine receptor gene from
Heliothis virescens (Lepidoptera: noctuidae)
Insect Biochem. Mol. Biol. 30 (4), 335-347 (2000)
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Puentes, R. and Windass, J.D.
Direct Submission
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